



Dexmedetomidine infusion for sedation of a patient with multiple system atrophy during spinal anesthesia

Satoko Kinouchi, MD^{1*}, Yoshie Onishi, MD¹,
Luna Nishihara, MD¹, Toshiki Okada, MD¹

ABSTRACT

¹Department of Anesthesiology, Minoh City Hospital, 5-7-1 Kayano, Mino-shi, Osaka, (Japan)

Correspondence:

Satoko Kinouchi, Department of Anesthesiology, Minoh City Hospital, 5-7-1 Kayano, Mino-shi, Osaka, 562-0014, (Japan); Telephone: 81-72-728-2001; Fax: 81-72-728-8475; E-mail: s.kinouchi@minoh-hp.jp

Received: 10 Mar 2018

Reviewed: 20 Mar 2018

Accepted: 5 Apr 2018

Multiple system atrophy (MSA) is a progressive neurodegenerative disorder characterized by extra-pyramidal symptoms, cerebellar ataxia, and autonomic dysfunction. The perioperative management of patients with MSA is challenging primarily due to autonomic dysfunction and vocal cord paralysis. Dexmedetomidine (DEX) is a highly selective α_2 -adrenoreceptor agonist that is used as a sedative, causing minimal depression of the respiratory function. We administered DEX continuously for sedation during spinal anesthesia for bipolar hip arthroplasty in a patient with MSA, without any cardiovascular or respiratory complications. This is the first case report of the efficacy of DEX for the sedation of a patient with MSA during spinal anesthesia.

Key words: Multiple system atrophy; Dexmedetomidine; Spinal anesthesia

Citation: Kinouchi S, Onishi Y, Nishihara L, Okada T. Dexmedetomidine infusion for sedation of a patient with multiple system atrophy during spinal anesthesia. *Anaesth Pain & Intensive Care* 2018;22(1):122-124

INTRODUCTION

The perioperative management of patients with MSA, a progressive neurodegenerative disorder,¹ is challenging, primarily due to autonomic dysfunction and vocal cord paralysis. Although dexmedetomidine (DEX) may be an excellent candidate as a sedative for patients with MSA during spinal anesthesia because of its minimal depression of the respiratory function, no case reports have been published on this topic in the medical literature.

CASE REPORT

A 67-year-old woman was scheduled to undergo bipolar hip arthroplasty for left femoral neck fracture. She had been suffered from gait disorder, dysarthria, incomplete voiding, and constipation for three years. On admission, she showed mild orthostatic hypotension: her blood pressure (BP) decreased

from 188/79 to 152/93 mmHg without any change in the heart rate (HR) on standing up from a supine position. Her symptoms, cerebellar dysfunction on neurological examination, and atrophy of the brain stem and cerebellum on magnetic resonance imaging (MRI) were all consistent with the diagnosis of MSA.

She was so anxious about surgery that she requested sedation during the operation. As we considered that general anesthesia might result in postoperative respiratory complications such as vocal cord paralysis and sleep apnea syndrome (SAS), we planned spinal anesthesia with sedation using DEX. In the operating theater, her BP, HR, and percutaneous arterial oxygen saturation (SpO₂) were 171/78 mmHg, 68 bpm, and 98%, respectively. Spinal anesthesia was performed with 11 mg of 0.5% hyperbaric bupivacaine, and anesthesia was achieved to T10, being sufficient to start the operation. Five minutes after the infusion

of DEX at a dose of 0.33 µg/kg/hr (loading dose), her Ramsay sedation score (RSS) had increased to 4, and the dose was subsequently decreased to 0.69 µg/kg/hr (maintenance dose). Because her BP and HR decreased from 172/78 to 110/57 mmHg and from 68 to 52 bpm, respectively, after the start of DEX infusion, the maintenance dose was titrated to 0.4 µg/kg/hr. The dose was further titrated to 0.28 µg/kg/hr thirty minutes after that, as her BP was 93/51 mmHg. Her hemodynamic state remained stable throughout the operation, without the necessity of any cardiovascular drugs.

She showed no signs of airway obstruction and her SpO₂ was maintained above 96% on room air during sedation. The operative time was 72 min. Five minutes after stopping DEX infusion when the operation finished, her RSS recovered to 2. She slept well, remembered nothing about the operation, and was satisfied with the procedure. She was discharged from the hospital 28 days after the surgery without any complications, and her MSA-associated symptoms remained unchanged.

DISCUSSION

This is the first published case report describing the safe and effective use of DEX for sedation of a patient with MSA during spinal anesthesia. MSA is a progressive neurodegenerative disorder characterized by extra-pyramidal symptoms, cerebellar ataxia, and autonomic dysfunction.²

MSA patients sometimes have respiratory complications such as vocal cord paralysis and SAS.³⁻⁶ In those with autonomic failure, their ventilatory responses to both hypoxia and hypercapnia have been reported to be markedly impaired,⁷ and they are also highly sensitive to tranquilizers and opioids.^{8,9} Those findings led us to select spinal rather than general anesthesia. Our patient wanted to be sedated during the operation since she was very anxious about being alert in the operating theater. Among the sedatives available, DEX has some notable advantages, which are markedly different from those of GABA-ergic sedatives such as propofol and benzodiazepines. An electroencephalogram (EEG) taken during DEX-induced sedation resembles that of normal sleep. As

a result, the muscle tone including that of respiratory muscles is well-preserved, and so its administration does not cause respiratory depression.¹⁰ The present patient showed no signs of airway obstruction and her SpO₂ was maintained above 96% in room air during sedation.

Autonomic dysfunction, one of the major symptoms of MSA, may lead to orthostatic hypotension due to a degenerated nucleus of the sympathetic system and/or unpredictable responses to vasopressors due to denervation hypersensitivity of the sympathetic system.¹¹⁻¹⁴ A case report demonstrated that patients with an already degenerated sympathetic system were unlikely to become more hypotensive on performing sympathetic blockade with spinal/epidural anesthesia.¹⁵ We selected spinal anesthesia with hyperbaric bupivacaine to easily control the level of anesthesia to as low as possible to perform the operation without pain.

DEX itself may cause cardiovascular problems because it can induce unpredictable changes such as hypertension, hypotension, and bradycardia, especially on high-dose infusion.^{16,17} Although DEX infusion for a patient with MSA might be challenging, its safe use was reported in an infant with familial dysautonomia.¹⁸ In our case, reduction of the DEX loading dose and titration of the maintenance dose were sufficient to maintain a stable BP and HR without requiring any cardiovascular drugs.

As a result, DEX infusion for sedation during spinal anesthesia may be a safe and effective option for patients with MSA. The accumulation of case reports is required to more definitively investigate the potential impact of α 2-adrenoreceptor agonists on patients with autonomic dysfunction.

Conflict of interest: Nil

Financial support: Nil

Authors' contribution:

SK — provided the anesthetic management and wrote the draft

YO, LN — reviewed the manuscript

TO — supervised the anesthetic management and reviewed the manuscript

REFERENCES

1. Gilman S, Wenning GK, Low PA, Brooks DJ, Mathias CJ, Trojanowski JQ, et al. Second consensus statement on the diagnosis of multiple system atrophy. *Neurology*. 2008;71(9):670-676. doi: 10.1212/01.wnl.0000324625.00404.15 [PubMed] [Free full text]
2. Wenning GK, Colosimo C, Geser F, Poewe W. Multiple system atrophy. *Lancet Neurol*. 2004;3:93-103. doi:10.1016/S1474-4422(03)00662-8. [PubMed]
3. Isozaki E, Naito A, Horiguchi S, Kawamura R, Hayashida T, Tanabe H. Early diagnosis and stage classification of vocal cord abductor paralysis in patients with multiple system atrophy. *J Neurol Neurosurg Psychiatry*. 1996;60(4):399-402. [PubMed] [Free full text]
4. Blumin JH, Berke GS. Bilateral vocal fold paresis and multiple system atrophy. *Arch Otolaryngol Head Neck Surg* 2002;128(12):1404-1407. [PubMed]
5. Oshima S, Sugihara K, Wakayama S. Aggravated sleep apnea after general anesthesia in a patient with Shy-Drager syndrome. *J Anesth*. 1994;8(4):484-486. doi: 10.1007/BF02514633 [PubMed]
6. Lim YS, Kennedy NJ. Multiple system atrophy as a cause of upper airway obstruction. *Anaesthesia*. 2007;62(11):1179-1182. [PubMed] Doi:10.1111/j.1365-2044.2007.05227.x
7. McNicholas WT, Rutherford R, Grossman R, Moldofsky H, Zamel N, Phillipson EA. Abnormal respiratory pattern generation during sleep in patients with autonomic dysfunction. *Am Rev Respir Dis*. 1983;128(3):429-433. [PubMed]
8. Sweeney BP, Jones S, Langford RM. Anaesthesia in dysautonomia: further complications. *Anaesthesia*. 1985;40(8):783-786. [PubMed]
9. Hanning CD. Obstructive sleep apnoea. *Br J Anaesth*. 1989;63(4):477-488. [PubMed]
10. Scott-Warren VL, Sebastian J. Dexmedetomidine: its use in intensive care medicine and anaesthesia. *BJA Education*. 2016;16(7):242-246. [Free full text]
11. Kluyskens Y, Bossaert L, Snoeck J, Martin JJ. Idiopathic orthostatic hypotension and the Shy and Drager syndrome: physiological studies in four cases-pathological report of one case. *Acta Cardiol*. 1977;32(5):317-335. [PubMed]
12. Bevan DR. Shy-Drager syndrome: a review and a description of the anaesthetic management. *Anaesthesia*. 1979;34(9):866-873. [PubMed]
13. Bannister R. Chronic autonomic failure with postural hypotension. *Lancet*. 1979;2(8139):404-406. [PubMed]
14. Stirt JA, Frantz RA, Gunz EF, Conolly ME. Anesthesia, catecholamines, and hemodynamics in autonomic dysfunction. *Anesth Analg*. 1982;61(8):701-704. [PubMed]
15. Cohen CA. Anesthetic management of a patient with the Shy-Drager Syndrome. *Anesthesiology*. 1971;35(1):95-97. [PubMed]
16. Lakhani PP, MacMillan LB, Guo TZ, McCool BA, Lovinger DM, Maze M, et al. Substitution of a mutant α 2-adrenergic receptor via 'hit and run' gene targeting reveals the role of this subtype in sedative, analgesic, and anesthetic-sparing responses in vivo. *Proc Natl Acad Sci U S A*. 1997;94(18):9950-9955. [PubMed] [Free full text]
17. Kamibayashi T, Maze M. Clinical uses of α 2-Adrenergic Agonists. *Anesthesiology*. 2000;93:1345-1349. [PubMed] [Free full text]
18. Abulhasan Y, Buu N, Frigon C. Perioperative use of dexmedetomidine in an infant with familial dysautonomia. *BJA*. 2009;103(3):413-415. doi: 10.1093/bja/aep178. [PubMed]

